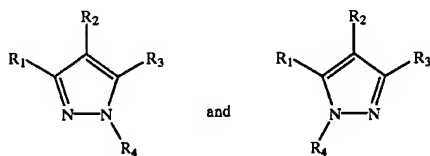


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- What is claimed is:
1. A compound having a formula selected from the group consisting of:



and their pharmaceutically acceptable salts, wherein:

- R₁ and R₃ are selected independently from the group consisting of optionally substituted hydroxyaryls and alkoxyaryls; R₂ is selected from the group consisting of hydrogen and optionally substituted loweralkyls and R₄ is selected from the group consisting of optionally substituted cycloalkyls.
2. The compound of claim 1, wherein R₁ and R₃ are selected independently from the group consisting of optionally substituted hydroxyaryls.
3. The compound of claim 1, wherein R₁ and R₃ are selected independently from the group consisting of optionally substituted alkoxyaryls.

4. The compound of claim 1, wherein at least one of R₁ and R₃ is substituted with at least one hydroxy or alkoxy group.
5. The compound of claim 1, wherein at least one of R₁ and R₃ is selected independently from the group consisting of optionally substituted phenyloxyloweralkyls.
6. The compound of claim 5, wherein at least one of R₁ and R₃ is substituted with a substituent selected from the group consisting of halogen, nitro, cyano, loweralkyl, haloloweralkyl, loweralkyloxy, haloloweralkyloxy, carboxy, loweralkyloxycarbonyl, aryloxycarbonyl, (cycloalkyl) oxycarbonyl, aralkyloxycarbonyl, heteroaryloxycarbonyl, heteroaralkyloxycarbonyl, (heterocycloalkyl) oxycarbonyl, loweralkylsulfinyl, loweralkylsulfonyl, loweralkylthio, arylthio, loweralkylcarbonyloxy, arylcarbonyloxy, aralkylcarbonyloxy, heteroarylcarbonyloxy, heteroaralkylcarbonyloxy, (cycloalkyl) carbonyloxy, alkylsulfonylamino, (heterocycloalkyl) carbonyloxy, aminocarbonyl, loweralkylaminocarbonyl, arylaminocarbonyl, aralkylaminocarbonyl, heteroarylaminocarbonyl, and heteroaralkylaminocarbonyl.
7. The compound of claim 6, wherein at least one of R₁ and R₃ is substituted with a substituent selected from the group consisting of halogen, nitro, cyano, loweralkyl, haloloweralkyl, loweralkyloxy, haloloweralkyloxy, carboxy, loweralkylthio, aminocarbonyl, and loweralkylsulfinyl.
8. The compound of claim 1, wherein R₂ is hydrogen.
9. The compound of claim 1, wherein R₂ is optionally substituted loweralkyl.
10. The compound of claim 1, wherein at least one of R₁ and R₃ is substituted with at least one hydroxy or thio group.
11. The compound of claim 1, wherein at least one of R₁ and R₃ is substituted with a substituent selected from the group consisting of halogen, loweralkyl, haloloweralkyl, loweralkyloxy, haloloweralkyloxy, carboxy, loweralkyloxycarbonyl, aryloxycarbonyl, (cycloalkyl) oxycarbonyl, aralkyloxycarbonyl, heteroaryloxycarbonyl, heteroaralkyloxycarbonyl, (heterocycloalkyl) oxycarbonyl, loweralkylsulfinyl, loweralkylsulfonyl, loweralkylthio, arylthio, loweralkylcarbonyloxy, arylcarbonyloxy, aralkylcarbonyloxy, heteroarylcarbonyloxy, heteroaralkylcarbonyloxy, (cycloalkyl) carbonyloxy, (heterocycloalkyl) carbonyloxy, aminocarbonyl, loweralkylaminocarbonyl, arylaminocarbonyl, aralkylaminocarbonyl, heteroarylaminocarbonyl, and heteroaralkylaminocarbonyl.
12. A composition for use in treating an estrogen receptor-mediated disorder in a mammal, comprising a therapeutically effective amount of a compound of claim 1 in a pharmaceutically effective carrier.

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